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Can we make cannabis safer?

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A changing cannabis climate

The cannabis landscape is rapidly changing. Following the Single Convention on Narcotic Drugs in 1961, possession, distribution and use were criminalised and cannabis-related arrests increased significantly in Europe and North America, particularly among younger and ethnic minority populations ¹. In subsequent decades cannabis use has waxed and waned in different countries, but there has been an overall trend towards greater use in most parts of the world ². This may be because prohibitive measures (scheduling substances or law enforcement) having little or no effect on use, abuse or substance related harms, as suggested by a recent systematic review ³. In the UK, the demand for treatment for cannabis-related problems increased by 56% in adults and 51% in those under 18 between the years 2005-06 and 2013-14 (Public Health England, 2014; 2015). Increased treatment-seeking for cannabis use problems has also been evident in the US ⁵, and in Europe, cannabis has become the primary illicit drug responsible for first-time entry to drug treatment, although in some countries this is mainly due to referrals from the criminal justice system ⁶.

Certain US states along with Uruguay have recently decided to allow cannabis to be sold for recreational purposes ⁷. Canada has recently proposed to legalise recreational use and several European countries have already lessened or dropped their criminal sanctions on cannabis possession and use ⁸. While it is likely that such moves will decrease the cost of crime associated with cannabis use, the effect they will have on cannabis consumption and the prevalence of cannabis associated harms is unclear.

In any event, moves toward legalization of medicinal or recreational cannabis are unlikely to decrease the number of people who use cannabis. However, they could facilitate measures to reduce population levels of harm – for example, by regulating cannabis potency and promoting safer (e.g. non-tobacco) routes of administration ⁹. It is therefore incumbent on

those concerned about cannabis-related harms to consider other ways in which the use of cannabis might be made safer.

Cannabis is becoming more potent

Cannabis Sativa L. contains at least 120 different compounds known as cannabinoids which are specific to the cannabis plant with more than 600 other compounds such as terpenoids and flavonoids ^{10,11}. The most abundant of the cannabinoids are delta-9-tetrahydrocannabinol (THC) and cannabidiol (CBD) which the plant produces in different ratios from the precursor cannabigerol (CBG), based on genes that code for certain enzymes (THCA-synthase and CBDA-synthase) ¹². Hence, increased THC content in cannabis will be at the cost of CBD content. Furthermore, preventing pollination of the female plant leads to a much more potent product as the plant converts the plants energy to producing more cannabinoids rather than seeds ¹³. This type of cannabis is referred to as *sinsemilla* which means “without seed” in Spanish, but is commonly termed ‘skunk’ in the UK.

Over the past 4 decades cannabis potency (its percentage of THC) has on average doubled worldwide ¹⁴ and CBD levels remaining low or absent in most cannabis preparations ^{15–18}. Systematic cross-breeding of cannabis plants by black-market growers has yielded a market dominance of high-THC plants. By 2008, roughly 80% of cannabis sold on the UK black market was high-potency cannabis ¹⁶, an increase from 55% in 2004-2005 ¹⁵ and only 15% in 1999-2002 ¹⁹. In the US, a recent report found that the increased THC levels in cannabis over the last two decades was due to a shift towards high-potency (*sinsemilla*) as opposed to regularly (outdoor) grown cannabis ¹⁸. Cannabis resin can be a more potent form of cannabis and has been found to reach potencies of over 60% THC (per weight) in the Netherlands ¹⁷, while in the UK resin (aka. hash) has low potency with equal quantities of THC (4%) and CBD (4%) ¹⁵. More recently, new extraction techniques are being used to produce extremely high-potency concentrates (e.g. Butane Hash Oil) up to 75% THC ²⁰. Even more concerning are synthetic cannabinoids (e.g. “Spice”) which typically act as full cannabinoid receptor agonists (THC is a partial agonist) and have been linked to severe adverse reactions including death ²¹.

The significance of CBD

The main possible adverse effects associated with cannabis use are dependence, cognitive and educational impairment and psychosis ²². Crucially, evidence suggests that the incidence of adverse consequences of cannabis use is associated with the amount of THC and CBD it contains. Encouragingly, it appears that increasing the CBD dose does not influence the pleasurable effects of THC. A recent study found that CBD given orally up to 800mg did not influence the pleasurable, reinforcing or intoxicating effects of a smoked cannabis cigarette (~4.4mg THC) ²³. Two studies of inhaled cannabis vapour, with combinations including 8mg THC, 8mg THC + 10mg CBD and 8mg THC + 16mg CBD, found no differences in ratings of 'stoned' ^{24,25}. Lastly, a study where participants smoked their own cannabis found no difference in 'stoned' ratings when comparing low to high CBD:THC ratio strains of cannabis ²⁶. Although research to date has found no impact of CBD on the pleasurable effects of THC, more research is warranted to confirm this.

Dependence

Approximately one in eleven people who try cannabis will become dependent in their lifetime ²⁷, although this risk is almost doubled if use starts in adolescence ²⁸ and between 25-50% among daily users ²⁹. However, not all varieties of cannabis have the same liability towards dependence. An online survey study of over 2514 cannabis users found that high-potency (high THC, low CBD) cannabis use was associated with a greater severity of cannabis dependence along with self-reported memory problems and paranoia, compared with use of lower potency cannabis ³⁰. However, high-potency cannabis was rated as the preferred type and the type producing the best "high". This is potentially important, as interventions aiming at switching users to a less potent variety may fail if the alternative lacks the pleasurable effects of high-potency cannabis.

A few case reports have found that CBD can reduce the symptoms associated with withdrawal from cannabis ^{31,32}, and two studies have found Sativex (CBD:THC ratio 1:1) to do the same ^{33,34} although it is not clear whether these effects were attributable to CBD, THC, or their interactive effects. Lastly, in an attentional bias task, cannabis containing

higher levels of CBD was found to reverse the heightened salience of cannabis and food cues compared to cannabis without CBD ²⁶. Therefore, varieties of cannabis with greater CBD content may reduce the likelihood of users developing dependence, whilst preserving the effects that users seek from cannabis.

Psychosis

High potency cannabis also carries a higher risk of psychosis as well as an earlier onset of the illness than low THC forms ^{35,36}. A recent case-control study of patients with first episode of psychosis found that daily use of high-THC, low CBD cannabis was associated with a 5-fold increase in the odds of psychosis but found no such increase among users of low potency hash (resin) ³⁷.

In a Dutch population study of 1,877 participants, those who used cannabis higher in CBD had experienced fewer lifetime psychotic-like experiences compared to those who preferred varieties with less CBD ³⁸. Some neuroimaging studies have also suggested that the MRI correlates of cannabis use may vary with exposure to CBD. One study found that regular use of high-potency cannabis was associated with alterations in the corpus callosum, an effect that was absent in hash users ³⁹. Two other MRI studies have reported that cannabis users had smaller hippocampal volumes than non-users, but that this was not evident in cannabis users with hair samples that were positive for CBD ^{40,41}. However, the extent to which cannabis use is associated with neuroanatomical changes remains controversial: studies that have matched participants on alcohol use, or have accounted for heritable and genetic risk, have failed to find an association between cannabis use and brain structure ⁴²⁻⁴⁴.

Cannabis use is particularly harmful for patients already suffering from a psychotic illness. A recent meta-analysis found that patients who continue to use cannabis are more likely to relapse than their non-using counterparts ⁴⁵. Moreover, continued use of high potency cannabis appears to be more harmful to patients with psychosis than use of cannabis with lower THC and a relatively high proportion of CBD ⁴⁶. Sadly, clinical interventions to reduce cannabis use in patients with psychosis have been largely unsuccessful ⁴⁷. A recent study in patients with a first episode of psychosis found that patients experience both the positive and negative effect of cannabis more intensely compared to healthy controls ⁴⁸. This may

explain their unwillingness to stop using in spite of the negative effect the drug has on them and the failure of interventions in this population. Another potential factor is that reducing substance use usually depend on the subject having motivation and insight, both of which are impaired in patients with psychotic disorders ⁴⁹.

Cognition and intelligence

There has been considerable debate regarding whether the use of cannabis may have lasting negative effects on memory functioning, intelligence and other aspects of cognition in the normal population ²². A recent study across three large samples found that cannabis use before the age of 17 was related to lower rates of high-school completion and degree attainment ⁵⁰. Similarly, a 1-year follow-up study of 1155 adolescents found that weekly cannabis use was related to poorer performance in GCSE Maths and English at age 16 (albeit less than tobacco), after controlling for confounders ⁵¹. However, a large twin study found that early school leaving was explained by shared environmental risk factors which increases the likelihood for both cannabis use and early school leaving ⁵². These findings suggest that although there is a link between cannabis use and poorer educational outcomes, these effects may be explained by other factors.

As for general intelligence, a 38-year follow-up study of 874 individuals found that those who had been dependent on cannabis for up to 20 years had experienced a drop in IQ of an average of 6 points ⁵³. However, this study was limited in terms of controlling for confounds and a relatively small number of the total study population in the affected group. Two recent large scale follow-up studies have found that cannabis use is not associated with reduced IQ when controlling for additional confounders and genetic factors ^{54,55}. These latter studies only followed participants up to the ages of 15 and 20 years and do not exclude the possibility that IQ might be affected if cannabis use and/or dependence is maintained for longer.

Although there remains a debate on whether or not cannabis use leads to long term cognitive impairments or educational problems following abstinence, it is however clear that ongoing regular cannabis use impairs cognition. A recent meta-analysis found that cannabis users perform significantly worse on memory tasks than non-users, with the greatest impairments being in prospective memory, visual recognition, immediate and

delayed recall ⁵⁶. The authors reported that regular use (defined as using between 4 to 20 times/month) or more was related to worse performance while no effect was seen in light users (4 times/month or less). However, no significant difference is seen in users compared to controls following 4 weeks of abstinence ⁵⁷. This is in line with PET imaging studies showing that downregulation of CB1 receptor densities can be reversed within 4 weeks of abstinence or earlier ^{58,59}.

Insights from cannabinoid experiments

While being fundamental to understand the population impact of a behaviour or exposure, epidemiological studies come with some clear limitations. When exploring different outcomes between cannabis users and non-users, one can never be certain that the observed effects (e.g. impaired cognition) are due to use itself or confounding variables. Hence, experimental studies such as randomised controlled trials (RCTs) are highly informative as the same person gets exposed to both experimental conditions or has a 50% chance of ending up in either condition. Unlike epidemiology, RCTs allow us to infer causality but with the caveat that it only allows conclusions relating to the acute effects of the drug and not from long-term exposure.

Experimental studies in healthy volunteers have shown that administration of high-dose THC can induce a range of psychological changes including transient psychotic symptoms ^{60–62}. They have also consistently found impairments in memory functioning in a dose response manner ^{60,63}. However, studies that have combined THC with CBD have found very different results. Co-administration of CBD significantly reduces THC-induced time estimation errors and psychological reactions ⁶⁴ while a subsequent study found 1mg/kg CBD significantly reduced the anxiogenic effects of 0.5mg/kg THC in healthy volunteers ⁶⁵. In a recent study of 140 cannabis users, those who tested positive for both THC and CBD in hair-samples experienced significantly fewer psychotic-like effects compared to those testing positive for THC only ⁶⁶. These results were later replicated in a study of recreational cannabis users ⁶⁷. The same group later administered inhaled THC (8mg) and CBD (16mg) to volunteers and found that CBD protected against the detrimental effects of THC on emotional processing ^{24, 68}. Furthermore, there is evidence that CBD reverses THC's illusory effects on depth perception

There is also evidence that CBD may reverse the negative impact of THC on cognitive performance. A naturalistic study of 134 cannabis users smoking their own cannabis found that participants using cannabis with higher CBD levels displayed no impairment on measures of immediate and delayed prose recall compared to when sober. By contrast, performance on these tasks was significantly impaired among those who used cannabis with equivalent THC but no CBD ⁶⁹. The same group explored memory functioning in 120 users while analysing hair-samples for presence of CBD. Participants who tested positive for CBD displayed significantly better performance ⁶⁷.

In our own research we found that pre-treatment with 600mg oral CBD before the administration of 1.5mg intravenous THC significantly inhibited paranoia, the occurrence of psychotic symptom and impairments to delayed recall in 48 healthy volunteers ⁷⁰. These effects were not attributable to changes in THC plasma levels in the CBD group. Rather, they appear to reflect opposite effects of THC and CBD on the brain regions that mediate psychotic and anxiety symptoms, and support cognitive processes like memory, emotional processing and response inhibition ^{71,72}.

Collectively, the available experimental evidence suggests that CBD, to a yet unknown extent, can mitigate the harmful effects of cannabis.

Making cannabis safer

It is vital, especially now that cannabis is becoming increasingly liberalised, that we explore alternative and innovative ways in which we can reduce and mitigate cannabis related harms.

Firstly, there should be more focus on the co-use of tobacco and cannabis and the additive harm this poses, especially since cannabis is frequently used together with tobacco, especially in Europe ⁷³. The use of other routes of administration such as smoke-free vaporisers have the potential of reducing the harmful effects the smoke poses to the user as well as reducing tobacco consumption ⁷⁴. This may be particularly important as tobacco increases the addictive potential of cannabis ^{75–77}. Furthermore, tobacco use may be an independent risk factor for psychosis ⁷⁸, as well as modifying the risk from cannabis use ⁷⁹. Studies of the impact of cannabis use on cognition have also found the relationship

weakened after controlling for use of tobacco ^{51,54,80}. However, controlled experimental studies on the interaction between cannabis and tobacco are currently lacking ⁸¹. Hence, it is vital that future studies address tobacco when evaluating the harms of cannabis.

Secondly, we need to better understand the harms posed by cannabis varieties with different THC potencies. Extremely potent cannabis concentrates (e.g. Butane Hash Oil) have gained popularity in the United States, where THC content is not currently regulated ⁸². Policies proposed by both Uruguay and the Netherlands of a 15% THC-cap could be beneficial, but are not yet based on a scientific understanding of harms posed by potencies above this limit ⁸³. Alternative approaches include taxation based on THC content, although further research is needed their impact (including use of more harmful, synthetic cannabinoids). Only very recently have studies differentiated between different types of cannabis based on THC content ^{30,37–39}, and most of these studies do not involve measures of THC and CBD content, but proxy measures of potency, such as analysis of police seizures or coffee shop cannabis, and a reliance on self-report measures. Although self-report measures are associated with THC and CBD content, these associations are modest and are weaker among infrequent users ^{84,85}. Future longitudinal studies may choose to collect cannabis cigarettes (joints) from their participants over the course of the study and collect information as to how often they would smoke such a cigarette. This will allow researchers to calculate a cumulative dose exposure of cannabinoids as well as tobacco – resulting in a far more accurate estimate of harm. This information could contribute to a cannabis use guideline, such as we currently have for alcohol.

Lastly, and perhaps most importantly, more knowledge is needed regarding the ratio between THC and CBD which reduces harm. As reviewed above, the available evidence suggests that CBD protects against many of the harms associated with THC; what is not known is the relative dose of CBD that is required to offset the negative effects of a given dose of THC. Experimental studies to date have been unable to establish this. Some have relied on the presence/absence of CBD in hair samples of participants ^{66,67}. Others have given CBD orally while administering inhaled or intravenous THC ^{23,70}. Cannabinoids are absorbed very differently orally as opposed to inhaled/intravenous (and result in differential metabolite profiles) and hence this does not allow an extrapolation of relative CBD:THC ratio ⁸⁶. Future experimental studies should therefore explore various CBD:THC ratios,

administered using the same route of administration, and using standardised measures of cognitive performance, psychopathology and liability for addiction. Once a “safer” CBD:THC ratio has been found, this may potentially be used as a harm-reduction strategy, where users (some also with a psychotic illness) experiencing negative effects from use of ‘skunk’-type cannabis can be encouraged to switch to the less harmful one. Encouragingly, studies to date have not found CBD to influence the rewarding or pleasurable effects of THC^{23–25}. This is crucial as any attempt to reduce the harms of cannabis would likely be ineffective if it also reduced the rewarding effects of cannabis.

Conclusion

In a rapidly changing political climate surrounding cannabis, the demand for effectively reducing cannabis related harms has never been greater, and more research (both experimental and observational) is urgently needed to inform policy decisions. Reducing the THC content of cannabis, smoke-free alternatives (vaporisers), and concurrent use of tobacco, may be effective harm reduction measures. However, increasing the content CBD may be especially promising as it can offset several harms of cannabis without compromising its rewarding effects.

Contributors

The article was conceptualised by AE and PM, and was written by AE. PM, RM and TF made additional comments, additional content recommendations and reviewed and corrected the manuscript

Declaration of interest

We declare no competing interests.

Search strategy and selection criteria

This was a narrative, critical review and was not carried out using standard search criteria and methods as for a systematic review. The articles used in producing this review obtained by means of searches through PubMed and Google Scholar, Google Scholar alerts for key terms ('cannabis', 'marijuana', 'cannabidiol') up to 07/10/16, reference lists in existing reviews and papers, and conference presentations.

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